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(54) METHODS OF CONTROLLING INSECTS

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(57) **ABSTRACT**

The present invention provides methods comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula (I): wherein $-B^1-B^2-B^3-$ is -C=N-O-, $-C=N-CH_2-$, $-C=CH_2-O-$ or $-N-CH_2-CH_2-$; L is a direct bond or methylene; A^1 and A^2 are C-H, or one of A^1 and A^2 is C-H and the other is N; X^1 is group X R^1 is $C_1\text{-}C_4\text{alkyl}$, $C_1\text{-}C_4\text{haloalkyl}$ or $C_3\text{-}C_6\text{cycloalkyl}$; R^2 is chlorodifluoromethyl or trifluoromethyl; each R^3 is independently bromo, chloro, fluoro or trifluoromethyl; R^4 is hydrogen, halogen, methyl, halomethyl or cyano; R^5 is hydrogen; or R^4 and R^5 together form a bridging 1,3-butadiene group; p is 2 or 3. Preferably the methods are for control of stemborer, leaffolder, hoppers, Gall midge, whorl maggot, Rice bugs, and/or Black bugs.

14 Claims, No Drawings

RELATED APPLICATION INFORMATION

This application is a 371 of International Application No. PCT/EP2013/066696, filed 09 Aug. 2013, which claims priority to EP Patent Application No. 12181766.2 filed 24 Aug. 2012 and EP Patent Application No. 12195026.5 filed 30 Nov. 2012 the contents of which are incorporated herein 10 by reference.

The present invention relates to a method of controlling insects, in particular insects that infest rice.

Compounds that are insecticidally, acaricidally, nemati- 15 cidally and/or moluscicidally active by antagnonism of the gamma-aminobutyric acid (GABA)-gated chloride channel, and which comprise a partially saturated heterocycle that is substituted by a haloalkyl substituent and one or two optionally substituted aromatic or heteroaromatic rings, represent 20 a new class of pesticides that are described for example in Ozoe et al. Biochemical and Biophysical Research Communications, 391 (2010) 744-749. Compounds from this class are broadly described in WO 2005/085216 (EP1731512), $_{25}$ WO 2007/123853, WO 2007/075459, WO 2009/002809, WO 2008/019760, WO 2008/122375, WO 2008/128711, WO 2009/097992, WO 2010/072781, WO 2010/072781, WO 2008/126665, WO 2007/125984, WO 2008/130651, JP 2008110971, JP 2008133273, JP 2009108046, WO 2009/ 022746, WO 2009/022746, WO 2010/032437, WO 2009/ 080250, WO 2010/020521, WO 2010/025998, WO 2010/ 020522, WO 2010/084067, WO 2010/086225, WO 2010/ 149506 and WO 2010/108733.

It has now surprisingly been found that particular insecticides from this new class of gamma-aminobutyric acid (GABA)-gated chloride channel antagonists (disclosed in e.g. WO2011/067272) are highly effective at controlling certain pests in rice.

These compounds therefore represent an important new solution for safeguarding crops of useful plants, particularly rice crops, from insects that infest rice crops, particularly where the insects are resistant to current methods.

In a first aspect the invention provides a method comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I

L is a direct bond or methylene;

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A¹ and A² are C—H, or one of A¹ and A² is C—H and the other is N: X1 is group X

 R^1 is C_1 - C_4 alkyl, C_1 - C_4 haloalkyl or C_3 - C_6 cycloalkyl; R² is chlorodifluoromethyl or trifluoromethyl; each R³ is independently bromo, chloro, fluoro or trifluoromethyl;

R⁴ is hydrogen, halogen, methyl, halomethyl or cyano; R⁵ is hydrogen;

or R⁴ and R⁵ together form a bridging 1,3-butadiene group; p is 2 or 3.

The method may be for controlling and/or preventing insects selected from the group consisting of stemborer, leaffolder, hoppers, Gall midge, whorl maggot, Rice bugs, and Black bugs.

In one embodiment the invention provides a method of controlling and/or preventing stemborer in rice comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I. In one embodiment the invention provides use of a compound of formula I for controlling and/or preventing stemborer, particularly in rice. The stemborer may be resistant to other 35 insecticides. Examples of stemborers include Chilo sp, Chilo suppressalis, Chilo polychrysus, Chilo auricilius, Scirpophaga spp., Scirpophaga incertulas, Scirpophaga innotata, Scirpophaga nivella Sesamia sp, Sesamia inferens. In a further aspect the invention provides a method for obtain-40 ing regulatory approval for the use of one or more of a compound of formula I to control stemborer, preferably in rice, comprising at least one step of referring to, submitting or relying on biological data showing that said active ingredient reduces insect pressure.

In one embodiment the invention provides a method of controlling and/or preventing leaffolder in rice comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I. In one embodiment the invention provides use of a compound 50 of formula I for controlling and/or preventing leaffolder, particularly in rice. The leaffolder may be resistant to other insecticides. Examples of leaffolders include Cnaphalocrocis spp., Cnaphalocrocis medinalis, Marasmia spp., Marasmia patnalis, Marasmia exigua. In a further aspect the invention provides a method for obtaining regulatory approval for the use of one or more of a compound of formula I to control leaffolder, preferably in rice, comprising at least one step of referring to, submitting or relying on biological data showing that said active ingredient reduces insect pressure.

In one embodiment the invention provides a method of controlling and/or preventing hoppers in rice comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I. In $-B^1-B^2-B^3$ is -C=N-O-, $-C=N-CH_2-$, 65 one embodiment the invention provides use of a compound of formula I for controlling and/or preventing hoppers, particularly in rice. The hoppers may be resistant to other

insecticides. Examples of Hoppers include Nephotettix spp., Nephotettix virescens, Nephotettix nigropictus, Nephotettix malayanus, Nephotettix cincticeps, Nilaparvata lugens, Sogatella furcifera. In a further aspect the invention provides a method for obtaining regulatory approval for the use 5 of one or more of a compound of formula I to control hoppers, preferably in rice, comprising at least one step of referring to, submitting or relying on biological data showing that said active ingredient reduces insect pressure.

In one embodiment the invention provides a method of controlling and/or preventing Gall midge in rice comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I. In one embodiment the invention provides use of a compound of formula I for controlling and/or preventing Gall midge, particularly in rice. The Gall midge may be resistant to other insecticides. Examples of Gall midge include Orseolia sp, Orseolia oryzae. In a further aspect the invention provides a method for obtaining regulatory approval for the use of one 20 or more of a compound of formula I to control Gall midge, preferably in rice, comprising at least one step of referring to, submitting or relying on biological data showing that said active ingredient reduces insect pressure.

In one embodiment the invention provides a method of 25 controlling and/or preventing whorl maggot in rice comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I. In one embodiment the invention provides use of a compound of formula I for controlling and/or preventing whorl maggot, 30 particularly in rice. The whorl maggot may be resistant to other insecticides. Examples of whorl maggots include Hydrellia sp, Hydrellia philippina. In a further aspect the invention provides a method for obtaining regulatory approval for the use of one or more of a compound of 35 formula I to control whorl maggots, preferably in rice, comprising at least one step of referring to, submitting or relying on biological data showing that said active ingredient reduces insect pressure.

In one embodiment the invention provides a method of 40 controlling and/or preventing Rice bugs in rice comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I. In one embodiment the invention provides use of a compound of formula I for controlling and/or preventing Rice bugs, 45 particularly in rice. The Rice bugs may be resistant to other insecticides. Examples of rice bugs include *Leptocorisa* sp. Leptocorisa oratorius, Leptocorisa chinensis, Leptocorisa acuta. In a further aspect the invention provides a method for obtaining regulatory approval for the use of one or more 50 of a compound of formula I to control Rice bugs, preferably in rice, comprising at least one step of referring to, submitting or relying on biological data showing that said active ingredient reduces insect pressure.

In one embodiment the invention provides a method of 55 controlling and/or preventing Black bugs in rice comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I. In one embodiment the invention provides use of a compound particularly in rice. The Black bugs may be resistant to other insecticides. Examples of Black bugs include Scotinophara sp, Scotinophara coarctata, Scotinophara lurida, Scotinophara latiuscula. In a further aspect the invention provides a method for obtaining regulatory approval for the use of one 65 or more of a compound of formula I to control Black bugs, preferably in rice, comprising at least one step of referring

to, submitting or relying on biological data showing that said active ingredient reduces insect pressure.

The compounds of formula (I) may exist in different geometric or optical isomers or tautomeric forms. This invention covers all such isomers and tautomers and mixtures thereof in all proportions as well as isotopic forms such as deuterated compounds. The invention also covers salts and N-oxides of the compounds of the invention.

Preferred substituent definitions are described below and may be combined in any combination, including with original definitions.

 $-B^1$ — B^2 — B^3 — is preferably —C—N—O—.

A¹ and A² are preferably C—H.

Preferably X¹ is 3,5-dichlorophenyl-, 3-chloro-4-fluorophenyl-, 3-fluoro-4-chlorophenyl-, 3,4-dichlorophenyl-, 3-chloro-4-bromophenyl-, 3,5-dichloro-4-fluorophenyl-, 3,4,5-trichlorophenyl-, 3,4,5-trifluorophenyl-, 3-chloro-5bromophenyl-, 3-chloro-5-fluorophenyl-, 3-chloro-5-(trifluoromethyl)phenyl-, 3,4-dichloro-5-(trifluoromethyl)phe-3,5-bis(trifluoromethyl)phenyl-, nvl-. 4-chloro-3.5-bis (trifluoromethyl)phenyl-, 3-(trifluoromethyl)phenyl-, more preferably 3-chloro-5-bromophenyl-, 3-chloro-5-(trifluoromethyl)phenyl-, 3,5-dichloro-4-fluorophenyl-, trichlorophenyl-, 3,5-bis(trifluoromethyl)phenyl-, 3-(trifluoromethyl)phenyl-, 3,5-dichloro-4-bromophenyl-, 3-bromo-5-(trifluoromethyl)phenyl-, 3,5-dibromophenyl-, or 3,4dichlorophenyl-, even more preferably 3,5-dichloro-phenyl, 3,5-dichloro-4-fluorophenyl-, 3,4,5-trichlorophenyl-, 3,5bis(trifluoromethyl)phenyl, most preferably R⁴ is 3,5-dichloro-phenyl.

R¹ is preferably methyl, ethyl, propyl, butyl, cyclopropyl, cyclobutyl, trifluoroethyl, difluoroethyl. Ethyl and trifluoroethyl are particularly preferred.

R² is preferably trifluoromethyl.

Preferably each R³ is independently chlorine or fluorine, most preferably chlorine.

R⁴ is preferably chloro or methyl, most preferably methyl. R⁵ is preferably hydrogen.

L is preferably a direct bond.

In one group of compounds $-B^1-B^2-B^3$ — is -C=N-O

In one group of compounds $-B^1-B^2-B^3$ — is -C=N--CH₂--

In one group of compounds $-B^1-B^2-B^3$ — is -C==CH₂--O-

In one group of compounds —B¹—B²—B³— is -N—CH₂—CH₂-

In one embodiment A¹ and A² are C—H; R² is trifluoromethyl, and R⁵ is hydrogen.

In one embodiment A¹ and A² are C—H, R² is trifluoromethyl, R⁵ is hydrogen and L is a direct bond.

In one embodiment $-B^1-B^2-B^3$ is -C=N-O. A¹ and A² are C—H, R² is trifluoromethyl, R⁵ is hydrogen and L is a direct bond.

In one embodiment $-B^1-B^2-B^3$ — is -C=N-O—. A¹ and A² are C—H, R² is trifluoromethyl, R⁴ is halogen or methyl, R⁵ is hydrogen and L is a direct bond.

In a preferred embodiment $-B^1-B^2-B^3$ — is $-C = N - O - A^1$ and A^2 are C - H, R^2 is trifluoromethyl, of formula I for controlling and/or preventing Black bugs, 60 R3 is chloro or fluoro, R4 is halogen or methyl, R5 is hydrogen and L is a direct bond.

> In a very preferred embodiment —B¹—B²—B³— is $-C = N - O - ... A^1$ and A^2 are C - H, R^2 is trifluoromethyl, R³ is chloro or fluoro, R⁴ is halogen or methyl, R⁵ is hydrogen and L is a direct bond, and the method is a method of controlling and/or preventing infestation of hoppers, preferably brown plant hopper (Nilaparvata lugens) or stem-

borer (in particular *Chilo* sp,) in a crop of useful plants, preferably rice, very preferably wherein the method is a method of controlling and/or preventing infestation of hoppers, preferably brown plant hopper (*Nilaparvata lugens*) in a crop of useful plants, preferably rice.

In one embodiment A¹ and A² are C—H, R² is trifluoromethyl, R⁴ is methyl, R⁵ is hydrogen, each R³ is chlorine, p is 2.

In one embodiment R^1 is C_1 - C_4 alkyl, e.g. methyl, ethyl or propyl, e.g. methyl or ethyl, e.g. ethyl.

In one embodiment X¹ is group Xa

In one embodiment R^1 is C_1 - C_5 alkyl, e.g. methyl, ethyl or propyl, e.g. methyl or ethyl, e.g. ethyl and X^1 is group Xa. 25

In one embodiment R¹ is methyl.

In one embodiment R¹ is ethyl.

In one embodiment R¹ is 2,2,2-trifluoroethyl.

In one embodiment R^1 is 2,2-difluoroethyl.

In one embodiment X¹ is 3,5-dichlorophenyl.

In one embodiment X¹ is 3,5-dichloro-4-fluorophenyl.

In one embodiment X^1 is 3,4,5-trichlorophenyl.

In one embodiment R^1 is methyl and X^1 is 3,5-dichlorophenyl.

In one embodiment R¹ is methyl and X¹ is 3,5-dichloro- ³⁵ 4-fluorophenyl.

In one embodiment R^1 is methyl and X^1 is 3,4,5-trichlorophenyl.

In one embodiment R^1 is ethyl and X^1 is 3,5-dichlorophenyl.

In one embodiment R^1 is ethyl and X^1 is 3,5-dichloro-4-fluorophenyl.

In one embodiment R^1 is ethyl and X^1 is 3,4,5-trichlorophenyl.

In one embodiment R^1 is 2,2,2-trifluoroethyl and X^1 is 3,5-dichlorophenyl.

In one embodiment R^1 is 2,2,2-trifluoroethyl and X^1 is 3,5-dichloro-4-fluorophenyl.

In one embodiment R^{Γ} is 2,2,2-trifluoroethyl and X^{Γ} is 50 3,4,5-trichlorophenyl.

In one embodiment R^1 is 2,2-diffuoroethyl and X^1 is 3,5-dichlorophenyl.

In one embodiment R^1 is 2,2-difluoroethyl and X^1 is 3,5-dichloro-4-fluorophenyl.

In one embodiment R^1 is 2,2-diffuoroethyl and X^1 is 3,4,5-trichlorophenyl.

In all embodiments of the invention the method is preferably a method of controlling and/or preventing infestation of hoppers, preferably brown plant hopper (*Nilaparvata* 60 *lugens*) or stemborer (in particular *Chilo* sp.) in a crop of useful plants, preferably rice. Method of controlling and/or preventing infestation of hoppers, preferably brown plant hopper (*Nilaparvata lugens*) in a crop of useful plants, preferably rice is most preferred.

Compounds of formula I may exist as compounds of formula I* or compounds of formula I**.

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$$\begin{array}{c} R^2 \\ H \\ R^3 \\ P \end{array}$$

Compounds of formula I** are more biologically active than compounds of formula I*. Compounds of formula I may be a mixture of compounds I* and I** in any ratio e.g. in a molar ratio of 1:99 to 99:1, e.g. 10:1 to 1:10, e.g. a substantially 50:50 molar ratio. Preferably the compound of formula I is a racemic mixture of the compounds of formula I** and I* or is enantiomerically enriched for the compound of formula I**. For example, when the compound of formula I is an enantiomerically enriched mixture of formula I**, the molar proportion of compound I** compared to the total amount of both enantiomers is for example greater than 50%, e.g. at least 55, 60, 65, 70, 75, 80, 85, 90, 95, 96, 97, 98, or at least 99%. In one embodiment the compound of formula I is a compound of formula I** in substantially pure form, e.g. it is provided substantially in the absence of the alternative enantiomer.

Compounds of formula I may also exist as compounds of $_{\rm 40}$ formula I' or compounds of formula I".

H
$$R^2$$
 R^3
 R^4
 R^5
 R^4
 R^4
 R^5
 R^4
 R^5
 R^4
 R^5
 R^4
 R^5
 R^6

(S=S stereochemistry, R=R stereochemistry)

Compounds of formula I" are often more biologically active than compounds of formula I'. The compound of formula I may be a mixture of compounds I' and I" in any ratio e.g. in a molar ratio of 1:99 to 99:1, e.g. 10:1 to 1:10,

e.g. a substantially 50:50 molar ratio. Preferably the compound of formula I is a racemic mixture of the compounds of formula I" and I' or is enantiomerically enriched for the compound of formula I". For example, when the compound of formula I is an enantiomerically enriched mixture of formula I", the molar proportion of compound I" compared to the total amount of both enantiomers is for example greater than 50%, e.g. at least 55, 60, 65, 70, 75, 80, 85, 90, 95, 96, 97, 98, or at least 99%. In a preferred embodiment the compound of formula I is a compound of formula I" in substantially pure form, e.g. it is provided substantially in the absence of the alternative enantiomer.

The above stereocentres give rise to four stereoisomers:

(I-iv)

In one embodiment the compound of formula I is a 55 mixture comprising compounds I-i, and I-iv, wherein the mixture is enriched for the compound of formula I-iv, e.g. the molar proportion of compound I-iv compared to the total amount of the four isomers is for example greater than 25%, e.g. at least 30, 35, 40, 50 55, 60, 65, 70, 75, 80, 85, 90, 95, 60 96, 97, 98, or at least 99%.

In another embodiment the compound of formula I is a mixture comprising compounds I-i, I-ii, I-iii and I-iv, wherein the molar amount of the compound of formula I-iv is greater than the molar amount of the compound of formula 65 I-i, and the molar amount of the compound I-ii, and the molar amount of the compound I-iii, in other

words, the compound of formula I-iv is the most abundant isomer in the mixture. For example the molar amount of compound of formula I-iv is at least 1, 2, 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 56, 70, 75, 80, 85, 90, or even at least 95% greater than the combined amount of the compound of formula I-iv and I-i, the combined amount of the compound of formula I-iv and I-ii, and the combined amount of the compound of formula I-iv and I-iii.

Although $B^1 - B^2 - B^3$ is shown above as C = N - O, the same applies in respect of the stereoisomers when $B^1 - B^2 - B^3$ is $-C = N - CH_2 - CH_2 - CH_2 - O$ and $-N - CH_2 -$

In one embodiment the compound of formula I-iv is the most abundant isomer and R^1 is C_1 - C_4 alkyl, e.g. methyl, ethyl or propyl, e.g. methyl or ethyl, e.g. ethyl.

In one embodiment the compound of formula I-iv is the most abundant isomer and R^1 is C_1 - C_4 alkyl, e.g. methyl, ethyl or propyl, e.g. methyl or ethyl, e.g. ethyl and X^1 is group Xa.

In one embodiment the compound of formula I-iv is the most abundant isomer and R¹ is methyl.

In one embodiment the compound of formula I-iv is the most abundant isomer and R^1 is ethyl.

In one embodiment the compound of formula I-iv is the most abundant isomer and R^1 is 2,2,2-trifluoroethyl.

In one embodiment the compound of formula I-iv is the most abundant isomer and R^1 is 2,2-difluoroethyl.

Preferred compounds of formula I are shown in the Tables below.

TABLE A

Compounds of formula (I-a)

Table A provides 78 compounds and mixtures of formula (I-a) wherein R^1 has the values listed in table X below. The symbols * and ** indicate the location of the chiral centres.

TABLE B

Compounds of formula (I-b)

$$F_3C$$
 $*$
 CI
 $*$

Table B provides 78 compounds and mixtures of formula (I-b) wherein R¹ has the values listed in table X below. The symbols * and ** indicate the location of the chiral centres.

TABLE C

Compounds of formula (I-c)		
F ₃ C N	(I-c)	5
F C_{l} H $**$ O N R^{l}		10

Table C provides 78 compounds and mixtures of formula $_{15}$ (I-c) wherein R^1 has the values listed in table X below. The symbols * and ** indicate the location of the chiral centres.

TABLE D

Compounds of formula (I-d)	
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	(I-d)

Table D provides 78 compounds and mixtures of formula (I-d) wherein R^1 has the values listed in table X below. The symbols * and ** indicate the location of the chiral centres.

TABLE E

Compounds of formula (I-e)

Table E provides 78 compounds and mixtures of formula 50 (I-e) wherein R^1 has the values listed in table X below. The symbols * and ** indicate the location of the chiral centres.

TABLE F

Compounds of formula (I-f)		55
F_3C $*$ Cl Cl Cl R	(I-f)	60
$O'' = R^{\perp}$		65

Table F provides 78 compounds and mixtures of formula (M) wherein R^1 has the values listed in table X below. The symbols * and ** indicate the location of the chiral centres.

Table X represents Table A when X is A, Table B when X is B, Table C when X is C, Table D when X is D, Table E when X is E, Table F when X is F.

10	Com-	Stereochemistry	Stereochemistry	
10	pound	at	at	
	numbers	*	**	R^1
	37.1	B 1 1.	B 1 1 1	.1 1
	X.1	Racemic mixture	Racemic mixture	ethyl-
	X.2	Racemic mixture	Racemic mixture	butyl-
	X.3	Racemic mixture	Racemic mixture	but-2-yl-
15	X.4	Racemic mixture	Racemic mixture	3-bromo-propyl-
	X.5	Racemic mixture	Racemic mixture	2,2,2-trifluoro-ethyl-
	X.6	Racemic mixture	Racemic mixture	3,3,3-trifluoro-propyl-
	X.7	Racemic mixture	Racemic mixture	cyclobutyl-
	X.8	Racemic mixture	Racemic mixture	methyl
	X.9	Racemic mixture	Racemic mixture	propyl
20	X.10	Racemic mixture	Racemic mixture	2,2-difluoro-ethyl-
	X.11	Racemic mixture	Racemic mixture	2-fluoro-ethyl-
	X.12	S	Racemic mixture	ethyl-
	X.13	S	Racemic mixture	butyl-
	X.14	S	Racemic mixture	but-2-yl-
	X.15	S	Racemic mixture	3-bromo-propyl-
	X.16	S	Racemic mixture	2,2,2-trifluoro-ethyl-
25	X.17	S	Racemic mixture	3,3,3-trifluoro-propyl-
	X.18	S	Racemic mixture	cyclobutyl-
	X.19	S	Racemic mixture	methyl
	X.20	S	Racemic mixture	propyl
	X.21	S	Racemic mixture	2,2-difluoro-ethyl-
	X.22	S	Racemic mixture	2-fluoro-ethyl-
30	X.22 X.23		Racemic mixture	•
		Racemic mixture		isopropyl
	X.24	Racemic mixture	Racemic mixture	cyclopropyl
	X.25	S	Racemic mixture	isopropyl
	X.26	S	Racemic mixture	cyclopropyl
	X.27	Racemic mixture	S	ethyl-
25	X.28	Racemic mixture	S	butyl-
35	X.29	Racemic mixture	S	but-2-yl-
	X.30	Racemic mixture	S	3-bromo-propyl-
	X.31	Racemic mixture	S	2,2,2-trifluoro-ethyl-
	X.32	Racemic mixture	S	3,3,3-trifluoro-propyl-
	X.33	Racemic mixture	S	cyclobutyl-
	X.34	Racemic mixture	S	methyl
40	X.34 X.35	Racemic mixture	S	•
				propyl
	X.36	Racemic mixture	S	2,2-difluoro-ethyl-
	X.37	Racemic mixture	S	2-fluoro-ethyl-
	X.38	S	S	ethyl-
	X.39	S	S	butyl-
45	X.40	S	S	but-2-yl-
	X.41	S	S	3-bromo-propyl-
	X.42	S	S	2,2,2-trifluoro-ethyl-
	X.43	S	S	3,3,3-trifluoro-propyl-
	X.44	S	S	cyclobutyl-
	X.45	S	S	methyl
50	X.46	S	S	propyl
50	X.47	S	S	2,2-difluoro-ethyl-
	X.48	S	S	
				2-fluoro-ethyl-
	X.49	Racemic mixture	S	isopropyl
	X.50	Racemic mixture	S	cyclopropyl
	X.51	S	S	isopropyl
55	X.52	S	S	cyclopropyl
	X.53	Racemic mixture	R	ethyl-
	X.54	Racemic mixture	R	butyl-
	X.55	Racemic mixture	R	but-2-yl-
	X.56	Racemic mixture	R	3-bromo-propyl-
	X.57	Racemic mixture	R	2,2,2-trifluoro-ethyl-
60	X.58	Racemic mixture	R	3,3,3-trifluoro-propyl-
50	X.59	Racemic mixture	R	cyclobutyl-
	X.60	Racemic mixture	R	methyl
	X.61	Racemic mixture	R	propyl
	X.62	Racemic mixture	R	2,2-difluoro-ethyl-
	X.63	Racemic mixture	R	2-fluoro-ethyl-
65	X.64	S	R	ethyl-
	X.65	S	R	butyl-

Com- pound numbers	Stereochemistry at *	Stereochemistry at **	R^1
X.66	S	R	but-2-yl-
X.67	S	R	3-bromo-propyl-
X.68	S	R	2,2,2-trifluoro-ethyl-
X.69	S	R	3,3,3-trifluoro-propyl-
X.70	S	R	cyclobutyl-
X.71	S	R	methyl
X.72	S	R	propyl
X.73	S	R	2,2-difluoro-ethyl-
X.74	S	R	2-fluoro-ethyl-
X.75	Racemic mixture	R	isopropyl
X.76	Racemic mixture	R	cyclopropyl
X.77	S	R	isopropyl
X.78	S	R	cyclopropyl

The compounds of the invention may be made by a variety of methods as shown in Schemes 1 to 3.

the ester directly to the amide by heating the ester and amine together in a thermal process. Suitable bases include pyridine, triethylamine, 4-(dimethylamino)-pyridine ("DMAP") or diisopropylethylamine (Hunig's base). Preferred solvents are N,N-dimethylacetamide, tetrahydrofuran, dioxane, 1,2-dimethoxyethane, ethyl acetate and toluene. The reaction is carried out at a temperature of from 0° C. to 100° C., preferably from 15° C. to 30° C., in particular at ambient temperature Amines of formula (III) are either known in the literature or can be prepared using methods known to a person skilled in the art.

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- Acid halides of formula (II), wherein R is Cl, F or Br, may be made from carboxylic acids of formula (II), wherein
 R is OH, under standard conditions, as described for example in WO2009/080250.
 - 3) Carboxylic acids of formula (II), wherein R is OH, may be formed from esters of formula (II), wherein R is C_1 - C_6 alkoxy as described for example in WO 2009/080250.

1) Compounds of formula (I), can be prepared by reacting 50 a compound of formula (II) wherein R is OH, C₁-C₆alkoxy or Cl, F or Br, with an amine of formula (III) as shown in Scheme 1. When R is OH such reactions are usually carried out in the presence of a coupling reagent, such as N,N'dicyclohexyl-carbodiimide ("DCC"), 1-ethyl-3-(3-dimeth-55 ylamino-propyl)carbodiimide hydrochloride ("EDC") or bis (2-oxo-3-oxazolidinyl)phosphonic chloride ("BOP-Cl"), in the presence of a base, and optionally in the presence of a nucleophilic catalyst, such as hydroxybenzotriazole ("HOBT"). When R is Cl, such reactions are usually carried out in the presence of a base, and optionally in the presence of a nucleophilic catalyst. Alternatively, it is possible to conduct the reaction in a biphasic system comprising an organic solvent, preferably ethyl acetate, and an aqueous 65 solvent, preferably a solution of sodium hydrogen carbonate. When R is C₁-C₆alkoxy it is sometimes possible to convert

- 4) Compounds of formula (I) can be prepared by reacting a compound of formula (IV) wherein X^B is a leaving group, for example a halogen, such as bromo, with carbon monoxide and an amine of formula (III), in the presence of a catalyst, such as palladium(II) acetate or bis-(triphenylphosphine)palladium(II) dichloride, optionally in the presence of a ligand, such as triphenylphosphine, and a base, such as sodium carbonate, pyridine, triethylamine, 4-(dimethylamino)-pyridine ("DMAP") or diisopropylethylamine (Hunig's base), in a solvent, such as water, N,N-dimethylformamide or tetrahydrofuran. The reaction is carried out at a temperature of from 50° C. to 200° C., preferably from 100° C. to 150° C. The reaction is carried out at a pressure of from 50 to 200 bar, preferably from 100 to 150 bar.
- 5) Compounds of formula (IV) wherein X^B is a leaving group, for example a halogen, such as bromo, can be made by a various of methods, for example as described in WO 2009/080250.

Scheme 2

$$X^B$$
 A^2
 A^1
 A^2
 A^3
 A^3
 A^3
 A^4
 A^3
 A^4
 A^3
 A^4
 A^3
 A^4
 A^3
 A^4
 A^4
 A^3
 A^4
 A^4

6) Alternatively, compounds of formula (I), can be prepared by various methods from an intermediate of formula $_{35}$ WO09/080250. An intermediate of formula (V) can be (V) as shown in Scheme 2 wherein X^B is a leaving group, for prepared for example from an intermediate of formula (VI) example a halogen, such as bromo, or X^B is cyano, formyl

or acetyl according to similar methods to those described in as described in the same reference.

$$\begin{array}{c} Scheme 3 \\ \hline \\ A^2 \\ A^1 \\ \hline \\ (Va) \\ (Va) \\ \hline \\ (Va) \\ (Va) \\ \hline \\ (Va) \\ \hline \\ (Va) \\ (Va) \\ \hline \\ (Va) \\ (Va) \\ \hline \\ (Va) \\$$

(I)

 $X^{C}-2$ 15

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7) Alternatively, compounds of formula (I) can be prepared by various methods from an intermediate of formula (VII) as shown in Scheme 3 wherein X^C is X^C -1 or X^C -2

$$X^{C-1}$$
 \mathbb{R}^2
 \mathbb{H}
 \mathbb{R}^3
 \mathbb{R}^3

$$\mathbb{R}^2$$
 \mathbb{H}
 \mathbb{H}
 \mathbb{R}^3

according to similar methods to those described in WO 25 2009/080250.

8) Compounds of formula (VII) wherein X^C is X^C is X^C -1 or X^C -2 can be prepared from a compound of formula (Va) from a compound of formula (VII) wherein X^C is CH_2 -halogen using similar methods to those described in WO 30 2009/080250.

9) Compounds of formula (VII) wherein X^{C} is CH_{2} -halogen, such as bromo or chloro, can be prepared by reacting a methyl ketone of formula (Va) with a halogenating agent, such as bromine or chlorine, in a solvent, such as 35 acetic acid, at a temperature of from 0° C. to 50° C., preferably from ambient temperature to 40° C.

Other methods for the preparation of compounds of formula I are described in PCT/EP2010/068605, which is incorporated herein by reference.

In one embodiment the invention provides a compound selected from Tables A to F for use against rice pests.

In one embodiment the invention provides a compound selected from Tables A to F for use against stemborer, particularly in rice.

Examples of stemborers include Chilo sp, Chilo suppressalis, Chilo polychrysus, Chilo auricilius, Scirpophaga spp., Scirpophaga incertulas, Scirpophaga innotata, Scirpophaga nivella Sesamia sp, Sesamia inferens.

In one embodiment the invention provides a compound 50 selected from Tables A to F for use against leaffolder, particularly in rice.

Examples of leaffolders include *Cnaphalocrocis* spp., *Cnaphalocrocis medinalis*, *Marasmia* spp., *Marasmia patnalis*, *Marasmia exigua*.

In one embodiment the invention provides a compound selected from Tables A to F for use against hoppers, particularly in rice.

Examples of Hoppers include Nephotettix spp., Nephotettix virescens, Nephotettix nigropictus, Nephotettix 60 malayanus, Nephotettix cincticeps, Nilaparvata lugens, Sogatella furcifera.

In one embodiment the invention provides a compound selected from Tables A to F for use against gallmidge, particularly in rice.

Examples of Gall midge include Orseolia sp, Orseolia

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In one embodiment the invention provides a compound selected from Tables A to F for use against whorl maggot, particularly in rice.

Examples of whorl maggots include *Hydrellia* sp, *Hydrel*⁵ *lia philippina*.

In one embodiment the invention provides a compound selected from Tables A to F for use against Rice bugs, particularly in rice.

Examples of rice bugs include *Leptocorisa* sp, *Leptocorisa oratorius*, *Leptocorisa chinensis*, *Leptocorisa acuta*.

In one embodiment the invention provides a compound selected from Tables A to F for use against Black bugs, particularly in rice.

Examples of Black bugs include Scotinophara sp, Scotinophara coarctata, Scotinophara lurida, Scotinophara latiuscula.

Use of the compounds of the invention against Brown plant hopper (*Nilaparvata lugens*) and/or stemborer (in particular *Chilo* sp.,) is particularly preferred. Use of the compounds of the invention against Brown plant hopper (*Nilaparvata lugens*) is most preferred.

The term "locus" of a useful plant as used herein is intended to embrace the place on which the useful plants are growing, where the plant propagation materials of the useful plants are sown or where the plant propagation materials of the useful plants will be placed into the soil. An example for such a locus is a field, on which crop plants are growing.

The term "plant propagation material" is understood to denote generative parts of a plant, such as seeds, which can be used for the multiplication of the latter, and vegetative material, such as cuttings or tubers, for example potatoes. There may be mentioned for example seeds (in the strict sense), roots, fruits, tubers, bulbs, rhizomes and parts of plants. Germinated plants and young plants which are to be transplanted after germination or after emergence from the soil, may also be mentioned. These young plants may be protected before transplantation by a total or partial treatment by immersion. Preferably "plant propagation material" is understood to denote seeds.

Application may be before infestation or when the pest is present. Application of the compounds of the invention can be performed according to any of the usual modes of application, e.g. foliar, drench, soil, in furrow etc. However, control of *Anthonomus grandis* s is usually achieved by foliar application, which is the preferred mode of application according to the invention.

Application of the compounds of the invention is preferably to a crop of cotton plants, the locus thereof or propagation material thereof.

The compounds of the invention may be applied to plant parts. Plant parts are to be understood as meaning all parts and organs of plants above and below the ground, such as shoot, leaf, flower and root, examples which may be mentioned being leaves, needles, stalks, stems, flowers, fruit bodies, fruits, seeds, roots, tubers and rhizomes. The plant parts also include harvested material, and vegetative and generative propagation material, for example cuttings, tubers, rhizomes, offshoots and seeds. Treatment according to the invention of the plants and plant parts with the active compounds can be carried out directly or by allowing the compounds to act on their surroundings, habitat or storage space by the customary treatment methods, for example by immersion, spraying, evaporation, fogging, scattering, painting on, injecting and, in the case of propagation material, in particular in the case of seed, also by applying one or more coats.

The compounds of the invention are suitable for use on any plant (preferably cotton plant), including those that have been genetically modified to be resistant to active ingredients such as herbicides, or to produce biologically active compounds that control infestation by plant pests.

The term "plant" as used herein includes seedlings, bushes and trees. Crops are to be understood as also including those crops which have been rendered tolerant to herbicides or classes of herbicides (e.g. ALS-, GS-, EPSPS-, PPO- and HPPD-inhibitors) by conventional methods of 10 breeding or by genetic engineering. An example of a crop that has been rendered tolerant to imidazolinones, e.g. imazamox, by conventional methods of breeding is Clearfield® summer rape (canola). Examples of crops that have been rendered tolerant to herbicides by genetic engineering 15 methods include e.g. glyphosate- and glufosinate-resistant maize varieties commercially available under the trade names RoundupReady® and LibertyLink®.

Compounds of formula I may be used on transgenic plants (including cultivars) obtained by genetic engineering methods and/or by conventional methods. These are understood as meaning plants having novel properties ("traits") which have been obtained by conventional breeding, by mutagenesis or by recombinant DNA techniques. Depending on the plant species or plant cultivars, their location and growth conditions (soils, climate, vegetation period, diet), the treatment according to the invention may also result in superadditive "synergistic") effects.

Thus, for example, reduced application rates and/or a widening of the activity spectrum and/or an increase in the 30 activity of the substances and compositions which can be used according to the invention, better plant growth, increased tolerance to high or low temperatures, increased tolerance to drought or to water or soil salt content, increased flowering performance, easier harvesting, accelerated maturation, higher harvest yields, higher quality and/or a higher nutritional value of the harvested products, better storage stability and/or processability of the harvested products are possible, which exceed the effects which were actually to be expected.

The preferred transgenic plants or plant cultivars which are to be treated according to the invention include all plants which, by virtue of the genetic modification, received genetic material which imparts particularly advantageous, useful traits to these plants. Examples of such traits are 45 better plant growth, increased tolerance to high or low temperatures, increased tolerance to drought or to water or soil salt content, increased flowering performance, easier harvesting, accelerated maturation, higher harvest yields, higher quality and/or a higher nutritional value of the 50 harvested products, better storage stability and/or processability of the harvested products.

Further and particularly emphasized examples of such traits are a better defence of the plants against animal and microbial pests, such as against insects, mites, phytopathosenic fungi, bacteria and/or viruses, and also increased tolerance of the plants to certain herbicidally active compounds.

Examples of transgenic plants which may be mentioned are the important crop plants, such as cereals (wheat, rice), 60 maize, soybean, potatoes, sugar beet, tomatoes, peas and other vegetable varieties, cotton, tobacco, oilseed rape and also fruit plants (with the fruits apples, pears, citrus fruits and grapes). Transgenic cotton is of particular interest.

Compounds of formula I may be used on transgenic plants 65 that are capable of producing one or more pesticidal proteins which confer upon the transgenic plant tolerance or resis-

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tance to harmful pests, e.g. insect pests, nematode pests and the like. Such pesticidal proteins include, without limitation, Cry proteins from Bacillus thuringiensis Cry1Ab, Cry1Ac, Cry1F, Cry2Ab, Cry2Ae, Cry3A, Cry3Bb, or Cry9C; engineered proteins such as modified Cry3A (U.S. Pat. No. 7,030,295) or Cry1A.105; or vegetative insecticidal proteins such as Vip1, Vip2 or Vip3. A full list of Bt Cry proteins and VIPs useful in the invention can be found on the worldwide web at Bacillus thuringiensis Toxin Nomenclature Database maintained by the University of Sussex (see also, Crickmore et al. (1998) Microbiol. Mol. Biol. Rev. 62:807-813). Other pesticidal proteins useful in the invention include proteins of bacteria colonizing nematodes, e.g. Photorhabdus spp. or Xenorhabdus spp.; toxins produced by animals, such as scorpion toxins, arachnid toxins, wasp toxins, or other insect-specific neurotoxins; toxins produced by fungi, such Streptomycetes toxins, plant lectins, such as pea or barley lectins; agglutinins; proteinase inhibitors, such as trypsin inhibitors, serine protease inhibitors, patatin, cystatin or papain inhibitors; ribosome-inactivating proteins (RIP), such as ricin, maize-RIP, abrin, luffin, saporin or bryodin; steroid metabolism enzymes, such as 3-hydroxysteroid oxidase, ecdysteroid-IDP-glycosyl-transferase, cholesterol oxidases, ecdysone inhibitors or HMG-CoA-reductase; ion channel blockers, such as blockers of sodium or calcium channels; juvenile hormone esterase; diuretic hormone receptors (helicokinin receptors); stilben synthase, bibenzyl synthase, chitinases or glucanases. Further examples of such pesticidal proteins or transgenic plants capable of synthesizing such proteins are disclosed, e.g., in EP-A 374753, WO 93/007278, WO 95/34656, EP-A 427529, EP-A 451878, WO 03/18810 and WO 03/52073. The methods for producing such transgenic plants are generally known to the person skilled in the art and some of which are commercially available such as Agrisure®CB (P1) (corn producing Cry1Ab), Agrisure®RW (P2) (corn producing mCry3A), Agrisure® Viptera (P3) (corn hybrids producing Vip3Aa); Agrisure300GT (P4) (corn hybrids producing Cry1Ab and mCry3A); YieldGard® (P5) (corn hybrids producing the Cry1Ab protein), YieldGard® Plus (P6) (corn hybrids producing Cry1Ab and Cry3Bb1), Genuity® SmartStax® (P7) (corn hybrids with Cry1A.105, Cry2Ab2, Cry1F, Cry34/35, Cry3Bb); Herculex® I (P8) (corn hybrids producing Cry1Fa) and Herculex®RW (P9) (corn hybrids producing Cry34Ab1, Cry35Ab1 and the enzyme Phosphinothricin-N-Acetyltransferase [PAT]); NuCOTN®33B (P10) (cotton cultivars producing Cry1Ac), Bollgard®I (P11) (cotton cultivars producing Cry1Ac), Bollgard®II (P12) (cotton cultivars producing Cry1Ac and Cry2Ab2) and VIPCOT® (P13) (cotton cultivars producing a Vip3Aa). Soybean Cyst Nematode resistance soybean (SCN®—Syngenta (P14)) and soybean with Aphid resistant trait (AMT® (P15)) are also of interest.

Further examples of such transgenic crops are:

1. Bt11 Maize from Syngenta Seeds SAS, Chemin de l'Hobit 27, F-31 790 St. Sauveur, France, registration number C/FR/96/05/10 (P16). Genetically modified *Zea mays* which has been rendered resistant to attack by the European corn borer (*Ostrinia nubilalis* and *Sesamia nonagrioides*) by transgenic expression of a truncated CryIA(b) toxin. Bt11 maize also transgenically expresses the enzyme PAT to achieve tolerance to the herbicide glufosinate ammonium.

2. Bt176 Maize from Syngenta Seeds SAS, Chemin de l'Hobit 27, F-31 790 St. Sauveur, France, registration number C/FR/96/05/10 (P17). Genetically modified *Zea mays* which has been rendered resistant to attack by the European corn borer (*Ostrinia nubilalis* and *Sesamia nonagrioides*) by

transgenic expression of a CryIA(b) toxin. Bt176 maize also transgenically expresses the enzyme PAT to achieve tolerance to the herbicide glufosinate ammonium.

3. MIR604 Maize from Syngenta Seeds SAS, Chemin de l'Hobit 27, F-31 790 St. Sauveur, France, registration num- 5 ber C/FR/96/05/10 (P18). Maize which has been rendered insect-resistant by transgenic expression of a modified CryIIIA toxin. This toxin is Cry3A055 modified by insertion of a cathepsin-D-protease recognition sequence. The preparation of such transgenic maize plants is described in WO 10 03/018810.

4. MON 863 Maize from Monsanto Europe S.A. 270-272 Avenue de Tervuren, B-1150 Brussels, Belgium, registration number C/DE/02/9 (P19). MON 863 expresses a CryIIIB (b1) toxin and has resistance to certain Coleoptera insects. 15

5. IPC 531 Cotton from Monsanto Europe S.A. 270-272 Avenue de Tervuren, B-1150 Brussels, Belgium, registration number C/ES/96/02. (P20)

6. 1507 Maize from Pioneer Overseas Corporation, Avenue Tedesco, 7 B-1160 Brussels, Belgium, registration 20 number C/NL/00/10. (P21) Genetically modified maize for the expression of the protein Cry1F for achieving resistance to certain Lepidoptera insects and of the PAT protein for achieving tolerance to the herbicide glufosinate ammonium.

7. NK603×MON 810 Maize from Monsanto Europe S.A. 25 270-272 Avenue de Tervuren, B-1150 Brussels, Belgium, registration number C/GB/02/M3/03. Consists of conventionally bred hybrid maize varieties by crossing the genetically modified varieties NK603 and MON 810. NK603× MON 810 Maize transgenically expresses the protein CP4 30 EPSPS, obtained from Agrobacterium sp. strain CP4, which imparts tolerance to the herbicide Roundup® (contains glyphosate), and also a CryIA(b) toxin obtained from Bacillus thuringiensis subsp. kurstaki which brings about tolerance to certain Lepidoptera, include the European corn 35

Further examples of transgenic plants, and of very high interest, are those carrying traits conferring resistance to 2.4 D (e.g. Enlist®) (e.g. WO 2011066384) (, glyphosate (e.g. Roundup Ready® (P24), Roundup Ready 2 Yield® (P25)), 40 sulfonylurea (e.g. STS®) (P26), glufosinate (e.g. Liberty Link® (P27), Ignite® (P28)), Dicamba (P29) (Monsanto), HPPD tolerance (P30) (e.g. isoxaflutole herbicide) (Bayer CropScience, Syngenta). Double or triple stacks of any of the traits described here are also of interest, including 45 glyphosate and sulfonyl-urea tolerance ((e.g. Optimum GAT®) (P31), plants stacked with STS® and Roundup Ready® (P32) or plants stacked with STS® and Roundup Ready 2 Yield® (P33)), dicamba and glyphosate tolerance (P34) (Monsanto). Of particular interest are soybean plants 50 carrying trains conferring resistance to 2.4 D (e.g. Enlist®), glyphosate (e.g. Roundup Ready®, Roundup Ready 2 Yield®), sulfonylurea (e.g. STS®), glufosinate (e.g. Liberty Link®, Ignite®), Dicamba (Monsanto) HPPD tolerance

Transgenic crops of insect-resistant plants are also described in BATS (Zentrum für Biosicherheit and Nachhaltigkeit, Zentrum BATS, Clarastrasse 13, 4058 Basel, Switzerland) Report 2003, (http://bats.ch).

Further examples of transgenic plants, and of very high 60 interest, are those carrying traits conferring resistance to 2.4 D (e.g. Enlist®) (e.g. WO 2011066384) (, glyphosate (e.g. Roundup Ready® (P24), Roundup Ready 2 Yield® (P25)), sulfonylurea (e.g. STS®) (P26), glufosinate (e.g. Liberty Link® (P27), Ignite® (P28)), Dicamba (P29) (Monsanto), 65 HPPD tolerance (P30) (e.g. isoxaflutole herbicide) (Bayer CropScience, Syngenta). Double or triple stacks of any of

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the traits described here are also of interest, including glyphosate and sulfonyl-urea tolerance ((e.g. Optimum GAT®) (P31), plants stacked with STS® and Roundup Ready® (P32) or plants stacked with STS® and Roundup Ready 2 Yield® (P33)), dicamba and glyphosate tolerance (P34) (Monsanto). Of particular interest are soybean plants carrying trains conferring resistance to 2.4 D (e.g. Enlist®), glyphosate (e.g. Roundup Ready®, Roundup Ready 2 Yield®), sulfonylurea (e.g. STS®), glufosinate (e.g. Liberty Link®, Ignite®), Dicamba (Monsanto) HPPD tolerance (e.g. isoxaflutole herbicide) (Bayer CropScience, Syngenta).

Examples of cotton transgenic events include MON 531/ 757/1076 (Bollgard I®-Monsanto), MON1445 (Roundup ready Cotton®-Monsanto), MON531 x MON1445 (Bollgard I+RR®—Monsanto), MON15985 (Genuity Bollgard II Cotton®-Monsanto), MON88913 (Genuity RR FLEX Cotton®-Monsanto), MON15985 x MON1445 (Genuity Bollgard II+RR FELX Cotton®-Monsanto), MON15983 x MON88913 (Genuity Bollgard II+RR FLEX Cotton®-Monsanto), MON15985 (FibreMax Bollgard II Cotton®-Monsanto), LL25 (FibreMax LL Cotton®-BCS Stoneville), GHB614 (FibreMax GlyTol Cotton®-BCS Stoneville), LL25 x MON15985 (FibreMax LL Bollgard II Cotton®-BCS Stoneville/Monsanto), GHB614 x LL25 (FibreMax LL GlyTol Cotton®-BCS Stoneville), GHB614 x LL25 x MON15985 (FibreMax RR GlyTol Bollgard II Cotton®-BCS Stoneville), MON88913 x MON15985 (FibreMax LL GlyTol Bollgard II Cotton®-Monsanto), MON88913 (Fibre-Max RR Flex Cotton®-Monsanto), GHB119+T304-40 (Twinlink®-BCS Stoneville), GHB119+T304-40 x LL25 x GHB614 (Twinlink LL GT®-BCS Stoneville), 3006-210-23×281-24-236 (PhytoGen Widestrike Insect Protection®-Dow), 3006-210-23×281-24-236 x MON88913 (PhytoGen Widestrike Insect Protection+RR FLEX-® Dow/Monsanto), 3006-210-23×281-24-236 x MON1445 ((PhytoGen Widestrike Insect Protection+RR®-Dow/Monsanto), MON1445 (PhytoGen Roundup Ready®-Monsanto), MON88913 (PhytoGen Roundup Ready FLEX®-Monsanto), COT102 x COT67B (Vipcot®-Syngenta), COT102 x COT67B x MON88913 (Vipcot RR FLEX®-Syngenta/Monsanto), 281-24-236 (Dow), 3006-210-23 (Dow), COT102 (Syngenta), COT67B (Syngenta), T304-40 (BCS Stoneville).

Examples of Soy transgenic events include MON87701 x MON89788 (Genuity Roundup ready 2 Yield Soybeans®-MON89788 (Roundup Ready2Yield®, Monsanto), RR2Y®-Monsanto), MON87708 (Monsanto), 40-3-2 (Roundup Ready®, RR1®-Monsanto), MON87701 (Monsanto), DAS-68416 (Enlist Weed Control System®-Dow), DP356043 (Optimum GAT®-Pioneer), A5547-127 (LibertyLink Soybean®-Bayercropscience), A2704-12 (Bayercropscience), GU262 (Bayercropscience), W62 W98 (Bayercropscience), CRV127 (Cultivance®-BASF/EM-BRAPA) SYHT0H2 (WO2012/082548)

Examples of Maize transgenic events include T25 (Lib-(e.g. isoxaflutole herbicide) (Bayer CropScience, Syngenta). 55 ertyLink®, LL®-Bayerscropscience), DHT-1 (Dow), TC1507 (Herculex I®-Dow), DAS59122-7 (Herculex TC1507+DAS59122-7-Herculex Xtra®-Dow), TC1507 x DAS-59122-7 x NK603 (Herculex Xtra+ RR®—Dow), TC1507 x DAS-59122 x MON88017 x MON89034 (Genuity Smartstax Corn®, Genuity Smartstax RIB Complete®-Monsanto/Dow), MON89034 x NK603 (Genuity VT double PRO®-Monsanto), MON89034+ MON88017 (Genuity VT Triple PRO®-Monsanto), NK603 (Roundup Ready 2®, RR2®-Monsanto), MON810 (Yield-Gard BT®, Yieldgard Cornborer®-Monsanto), MON810 x NK603 (YieldGard cornborer RR Corn 2®-Monasnto), MON810 x MON863 (YieldGard Plus®—Monsanto),

MON863 x MON810 x NK603 (YieldGard Plus+RR Corn2®/YieldGard RR Maize®—Monsanto), MON863 x NK603 (YieldGard Rotworm+RR Corn 2®-Monsanto), MON863 (YieldBard RW®-Monsanto), MON89034 (Yield-Gard RW®-Monsanto), MON88017 (YieldGard VT RW®—Monsanto), MON810+MON88017 (YieldGard VT Triple®-Monsanto), MON88017+MON89034 (YieldGard VT Triple Pro®-Monsanto), Bt11+MIR604+GA21 (Agrisure 3000®-Syngenta), Bt11+TC1507+MIR604+5307+ GA21 (Syngenta), Bt11+TC1507+MIR604+DAS59122+ GA21 (Agrisure 3122®-Syngenta), BT11 (Agrisure CB®-Syngenta), GA21-(Agrisure GT®-Syngenta), MIR604 (Agrisure RW®-Syngenta), Bt11+MIR162 (Agrisure TL VIP®-Syngenta), BT11+MIR162+GA21 (Agrisure Viptra 31100-Syngenta), BT11+MIR162+MIR604 (AgrisureTM Event3272+BT11+MIR604+GA21 3100®—Syngenta), (Syngenta), BT11+MIR1692+MIR604+GA21 (Agrisure Viptera 3111®-Syngenta), BT11+MIR162+TC1507+GA21 (Agrisure Viptera 3220®—Syngenta), BT11+MIR162+ TC1507+MIR604+5307+GA21 (Agrisure Viptera 3222®- 20 Syngenta), MIR162 (Syngenta), BT11+GA21+MIR162+ MIR604+5307 (Syngenta), 5307 (Syngenta).

Herbicide-resistant plants (plants bred in a conventional manner for herbicide tolerance) which may be mentioned include the varieties sold under the name Clearfield®) (for 25 e) Organic tin compounds, such as cyhexatin, fenbutatin example maize).

These statements also apply to plant cultivars having these genetic traits or genetic traits still to be developed, which plant cultivars will be developed and/or marketed in the future.

The compounds of the invention are suitable for use on any cotton plant, including those that have been genetically modified to be resistant to active ingredients such as herbicides, or to produce biologically active compounds that control infestation by plant pests, e.g. BT cotton.

A compound of the invention may be used in mixtures with fertilizers (for example nitrogen-, potassium- or phosphorus-containing fertilizers). Suitable formulation types include granules of fertilizer. The mixtures preferably con-

The invention therefore also provides a fertilizer composition comprising a fertilizer and a compound of the invention.

The compositions of this invention may contain other compounds having biological activity, for example micro- 45 nutrients or compounds having fungicidal activity or which possess plant growth regulating, herbicidal, insecticidal, nematicidal or acaricidal activity.

The compositions of this invention may contain other compounds having biological activity, for example micro- 50 nutrients or compounds having fungicidal activity or which possess plant growth regulating, herbicidal, insecticidal, nematicidal or acaricidal activity.

The compound of formula (I) may be the sole active ingredient of the composition or it may be admixed with one 55 or more additional active ingredients such as a pesticide, e.g. a insecticide, fungicide or herbicide, or a synergist or plant growth regulator where appropriate. An additional active ingredient may provide a composition having a broader spectrum of activity or increased persistence at a locus; synergize the activity or complement the activity (for example by increasing the speed of effect or overcoming repellency) of the compound of formula (I); or help to overcome or prevent the development of resistance to individual components. The particular additional active ingre- 65 dient will depend upon the intended utility of the composition. Examples of suitable pesticides include the following:

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- a) Pyrethroids, such as permethrin, cypermethrin, fenvalerate, esfenvalerate, deltamethrin, cyhalothrin (in particular lambda-cyhalothrin and gamma cyhalothrin), bifenthrin, fenpropathrin, cyfluthrin, tefluthrin, fish safe pyrethroids (for example ethofenprox), natural pyrethrin, tetramethrin, S-bioallethrin, fenfluthrin, prallethrin, acrinathirin, etofenprox or 5-benzyl-3-furylmethyl-(E)-(1R,3S)-2,2dimethyl-3-(2-oxothiolan-3-ylidenemethyl)cyclopropane carboxylate;
- 10 b) Organophosphates, such as profenofos, sulprofos, acephate, methyl parathion, azinphos-methyl, demeton-smethyl, heptenophos, thiometon, fenamiphos, monocrotophos, profenofos, triazophos, methamidophos, dimephosphamidon, thoate. malathion, chlorpyrifos, phosalone, terbufos, fensulfothion, fonofos, phorate, phoxim, pirimiphos-methyl, pirimiphos-ethyl, fenitrothion, fosthiazate or diazinon;
 - c) Carbamates (including aryl carbamates), such as pirimicarb, triazamate, cloethocarb, carbofuran, furathiocarb, ethiofencarb, aldicarb, thiofurox, carbosulfan, bendiocarb, fenobucarb, propoxur, methomyl or oxamyl;
 - d) Benzoyl ureas, such as diflubenzuron, triflumuron, hexaflumuron, flufenoxuron, diafenthiuron, lufeneron, novaluron, noviflumuron or chlorfluazuron;
 - oxide or azocyclotin;
 - f) Pyrazoles, such as tebufenpyrad, tolfenpyrad, ethiprole, pyriprole, fipronil, and fenpyroximate;
 - g) Macrolides, such as avermectins or milbemycins, for example abamectin, emamectin benzoate, ivermectin, milbemycin, spinosad, azadirachtin, milbemectin, lepimectin or spinetoram;
 - h) Hormones or pheromones;
 - i) Organochlorine compounds, such as endosulfan (in particular alpha-endosulfan), benzene hexachloride, DDT, chlordane or dieldrin:
 - j) Amidines, such as chlordimeform or amitraz;
 - k) Fumigant agents, such as chloropicrin, dichloropropane, methyl bromide or metam;
- tain up to 25% by weight of the compound of the invention. 40 1) Neonicotinoid compounds, such as imidacloprid, thiacloprid, acetamiprid, nitenpyram, dinotefuran, thiamethoxam, clothianidin, or nithiazine;
 - m) Diacylhydrazines, such as tebufenozide, chromafenozide or methoxyfenozide;
 - n) Diphenyl ethers, such as diofenolan or pyriproxifen;
 - o) Ureas such as Indoxacarb or metaflumizone;
 - p) Ketoenols, such as Spirotetramat, spirodiclofen or spiromesifen;
 - q) Diamides, such as flubendiamide, chlorantraniliprole (Rynaxypyr®) or cyantraniliprole;
 - r) Essential oils such as Bugoil®—(PlantImpact); or
 - s) a comopund selected from buprofezine, flonicamid, acequinocyl, bifenazate, cyenopyrafen, cyflumetofen, etoxazole, flometoquin, fluacrypyrim, fluensulfone, flufenerim, flupyradifuone, harpin, iodomethane, dodecadienol, pyridaben, pyridalyl, pyrimidifen, flupyradifurone, 4-[(6-Chloro-pyridin-3-ylmethyl)-(2,2-difluoro-ethyl)-amino]-5H-furan-2-one (DE 102006015467), CAS: 915972-17-7 (WO 2006129714; WO2011/147953; WO2011/147952), CAS: 26914-55-8 (WO 2007020986), chlorfenapyr, pymetrozine, sulfoxaflor and pyrifluqinazon.

In addition to the major chemical classes of pesticide listed above, other pesticides having particular targets may be employed in the composition, if appropriate for the intended utility of the composition. For instance, selective insecticides for particular crops, for example stemborer specific insecticides (such as cartap) or hopper specific

acid

amide.

insecticides (such as buprofezin) for use in rice may be employed. Alternatively insecticides or acaricides specific for particular insect species/stages may also be included in the compositions (for example acaricidal ovo-larvicides, such as clofentezine, flubenzimine, hexythiazox or tetradifon; acaricidal motilicides, such as dicofol or propargite; acaricides, such as bromopropylate or chlorobenzilate; or growth regulators, such as hydramethylnon, cyromazine, methoprene, chlorfluazuron or diflubenzuron).

Examples of fungicidal compounds which may be 10 included in the composition of the invention are (E)-Nmethyl-2-[2-(2,5-dimethylphenoxymethyl)phenyl]-2methoxy-iminoacetamide (SSF-129), 4-bromo-2-cyano-N, N-dimethyl-6-trifluoromethylbenzimidazole-1-sulfonamide, α -[N-(3-chloro-2,6-xylyl)-2-methoxyacetamido]- γ - 15 butyrolactone, 4-chloro-2-cyano-N,N-dimethyl-5-ptolylimidazole-1-sulfonamide cyamidazosulfamid), 3-5-dichloro-N-(3-chloro-1-ethyl-1methyl-2-oxopropyl)-4-methylbenzamide (RH-7281, zoxamide), N-allyl-4,5,-dimethyl-2-trimethylsilylthiophene-3- 20 (MON65500), N-(1-cyano-1,2dimethylpropyl)-2-(2,4-dichlorophenoxy)propionamide (AC382042), N-(2-methoxy-5-pyridyl)-cyclopropane carboxamide, acibenzolar (CGA245704) (e.g. acibenzolar-Smethyl), alanycarb, aldimorph, anilazine, azaconazole, 25 azoxystrobin, benalaxyl, benomyl, benthiavalicarb, biloxazol, bitertanol, bixafen, blasticidin S, boscalid, bromuconazole, bupirimate, captafol, captan, carbendazim, carbendazim chlorhydrate, carboxin, carpropamid, carvone, CGA41396, CGA41397, chinomethionate, chlorothalonil, 30 chlorozolinate, clozylacon, copper containing compounds such as copper oxychloride, copper oxyquinolate, copper sulfate, copper tallate and Bordeaux mixture, cyclufenamid, cymoxanil, cyproconazole, cyprodinil, debacarb, di-2pyridyl disulfide 1,1'-dioxide, dichlofluanid, diclomezine, 35 dicloran, diethofencarb, difenoconazole, difenzoquat, diflumetorim, O,O-di-iso-propyl-S-benzyl thiophosphate, dimefluazole, dimetconazole, dimethomorph, dimethirimol, diniconazole, dinocap, dithianon, dodecyl dimethyl ammonium chloride, dodemorph, dodine, doguadine, edifenphos, epoxi- 40 ethyl-(Z)—N-benzyl-N-([methyl conazole, ethirimol, (methyl-thioethylideneamino-oxycarbonyl)amino thio)-βalaninate, etridiazole, famoxadone, fenamidone (RPA407213), fenarimol, fenbuconazole, fenfuram, fenhexamid (KBR2738), fenpiclonil, fenpropidin, fenpropimorph, 45 fentin acetate, fentin hydroxide, ferbam, ferimzone, fluazinam, fludioxonil, flumetover, fluopyram, fluoxastrobin, fluoroimide, fluquinconazole, flusilazole, flutolanil, flutriafol, fluxapyroxad, folpet, fuberidazole, furalaxyl, furametpyr, guazatine, hexaconazole, hydroxyisoxazole, hymexazole, 50 imazalil, imibenconazole, iminoctadine, iminoctadine triacetate, ipconazole, iprobenfos, iprodione, iprovalicarb (SZX0722), isopropanyl butyl carbamate, isoprothiolane, isopyrazam, kasugamycin, kresoxim-methyl, LY186054, LY211795, LY248908, mancozeb, mandipropamid, maneb, 55 mefenoxam, metalaxyl, mepanipyrim, mepronil, metalaxyl, metconazole, metiram, metiram-zinc, metominostrobin, myclobutanil, neoasozin, nickel dimethyldithiocarbamate, nitrothal-isopropyl, nuarimol, ofurace, organomercury compounds, oxadixyl, oxasulfuron, oxolinic acid, oxpoconazole, 60 oxycarboxin, pefurazoate, penconazole, pencycuron, penflufen, penthiopyrad, phenazin oxide, phosetyl-Al, phosphorus acids, phthalide, picoxystrobin (ZA1963), polyoxinD, polyram, probenazole, prochloraz, procymidone, propamocarb, propiconazole, propineb, propionic acid, prothioconazole, pyrazophos, pyrifenox, pyrimethanil, pyraclostrobin, pyroquilon, pyroxyfur, pyrrolnitrin, quaternary ammonium

compounds, quinomethionate, quinoxyfen, quintozene, sedaxane, sipconazole (F-155), sodium pentachlorophenate, spiroxamine, streptomycin, sulfur, tebuconazole, tecloftalam, tecnazene, tetraconazole, thiabendazole, thifluzamid, 2-(thiocyanomethylthio)benzothiazole, thiophanate-methyl, thiram, timibenconazole, tolclofos-methyl, tolylfluanid, triadimefon, triadimenol, triazbutil, triazoxide, tricyclazole, tridemorph, trifloxystrobin (CGA279202), triforine, triflumizole, triticonazole, validamycin A, vapam, vinclozolin, zineb and ziram, N-[9-(dichloromethylene)-1,2,3,4-tetrahydro-1,4-methanonaphthalen-5-yl]-3-(difluoromethyl)-1methyl-1H-pyrazole-4-carboxamide[1072957-71-1], 1-methyl-3-difluoromethyl-1H-pyrazole-4-carboxylic acid (2-dichloromethylene-3-ethyl-1-methyl-indan-4-yl)-amide, and 1-methyl-3-difluoromethyl-4H-pyrazole-4-carboxylic

[2-(2,4-dichloro-phenyl)-2-methoxy-1-methyl-ethyl]-

In addition, biological agents may be included in the composition of the invention e.g. *Baciullus* species such as *Bacillus firmus*, *Bacillus cereus*, *Bacillus subtilis*, and *Pasteuria* species such as *Pasteuria penetrans* and *Pasteuria nishizawae*. A suitable *Bacillus firmus* strain is strain CNCM 1-1582 which is commercially available as BioNemTM. A suitable *Bacillus cereus* strain is strain CNCM 1-1562. Of both *Bacillus* strains more details can be found in U.S. Pat. No. 6,406,690. Other biological organisms that may be included in the compositions of the invention are bacteria such as *Streptomyces* spp. such as *S. avermitilis*, and fungi such as *Pochonia* spp. such as *P. chlamydosporia*. Also of interest are *Metarhizium* spp. such as *M. anisopliae*; *Pochonia* spp. such as *P. chlamydosporia*.

The compounds of the invention may be mixed with soil, peat or other rooting media for the protection of plants against seed-borne, soil-borne or foliar fungal diseases.

Examples of suitable synergists for use in the compositions include piperonyl butoxide, sesamex, safroxan and dodecyl imidazole.

Suitable herbicides and plant-growth regulators for inclusion in the compositions will depend upon the intended target and the effect required.

An example of a rice selective herbicide which may be included is propanil. An example of a plant growth regulator for use in cotton is PIXTM

Some mixtures may comprise active ingredients which have significantly different physical, chemical or biological properties such that they do not easily lend themselves to the same conventional formulation type. In these circumstances other formulation types may be prepared. For example, where one active ingredient is a water insoluble solid and the other a water insoluble liquid, it may nevertheless be possible to disperse each active ingredient in the same continuous aqueous phase by dispersing the solid active ingredient as a suspension (using a preparation analogous to that of an SC) but dispersing the liquid active ingredient as an emulsion (using a preparation analogous to that of an EW). The resultant composition is a suspoemulsion (SE) formulation.

Unless otherwise stated the weight ratio of the compound of I with an additional active ingredient may generally be between 1000:1 and 1:1000. In other embodiments that weight ratio of the compound of formula I to the additional active ingredient may be between 500:1 to 1:500, for example between 100:1 to 1:100, for example between 1:50 to 50:1, for example 1:20 to 20:1, for example 1:10 to 10:1, for example 1:5 to 5:1, for example 1:1, 1:2, 1:3, 1:4, 1:5, 2:1, 3:1, 4:1, or 5:1.

Mixtures with pyrethroids, in particular pymetrozine, are of particular interest for the present invention.

Compositions of the invention include those prepared by premixing prior to application, e.g. as a readymix or tankmix, or by simultaneous application or sequential application to the plant.

In order to apply a compounds of the invention as an 5 insecticide, acaricide, nematicide or molluscicide to a pest, a locus of pest, or to a plant susceptible to attack by a pest, compounds of the invention is usually formulated into a composition which includes, in addition to the compound of the invention, a suitable inert diluent or carrier and, option- 10 ally, a surface active agent (SFA). SFAs are chemicals which are able to modify the properties of an interface (for example, liquid/solid, liquid/air or liquid/liquid interfaces) by lowering the interfacial tension and thereby leading to changes in other properties (for example dispersion, emul- 15 sification and wetting). It is preferred that all compositions (both solid and liquid formulations) comprise, by weight, 0.0001 to 95%, more preferably 1 to 85%, for example 5 to 60%, of a compound of the invention. The composition is generally used for the control of pests such that a compound 20 of the invention is applied at a rate of from 0.1 g to 10 kg per hectare, preferably from 1 g to 6 kg per hectare, more preferably from 1 g to 1 kg per hectare.

In one embodiment the compounds of the invention are used for pest control on cotton at 1:500 g/ha, for example 25 10-70 g/ha. However, it should be noted that due to the very damaging effect of the *Anthonomus grandis* (quantity and quality on yield), sprays are often very intense and at very low threshold levels and can be down to almost zero tolerance.

When used in a seed dressing, a compound of the invention is used at a rate of 0.0001 g to 10 g (for example 0.001 g or 0.05 g), preferably 0.005 g to 10 g, more preferably 0.005 g to 4 g, per kilogram of seed.

Compositions comprising a compound of the invention 35 can be chosen from a number of formulation types, including dustable powders (DP), soluble powders (SP), water soluble granules (SG), water dispersible granules (WG), wettable powders (WP), granules (GR) (slow or fast release), soluble concentrates (SL), oil miscible liquids 40 (OL), ultra low volume liquids (UL), emulsifiable concentrates (EC), dispersible concentrates (DC), emulsions (both oil in water (EW) and water in oil (EO)), micro-emulsions (ME), suspension concentrates (SC), aerosols, fogging/smoke formulations, capsule suspensions (CS) and seed 45 treatment formulations. The formulation type chosen in any instance will depend upon the particular purpose envisaged and the physical, chemical and biological properties of the compound of the invention.

Dustable powders (DP) may be prepared by mixing a 50 compound of the invention with one or more solid diluents (for example natural clays, kaolin, pyrophyllite, bentonite, alumina, montmorillonite, kieselguhr, chalk, diatomaceous earths, calcium phosphates, calcium and magnesium carbonates, sulfur, lime, flours, talc and other organic and 55 inorganic solid carriers) and mechanically grinding the mixture to a fine powder.

Soluble powders (SP) may be prepared by mixing a compound of the invention with one or more water-soluble inorganic salts (such as sodium bicarbonate, sodium carbonate or magnesium sulfate) or one or more water-soluble organic solids (such as a polysaccharide) and, optionally, one or more wetting agents, one or more dispersing agents or a mixture of said agents to improve water dispersibility/solubility. The mixture is then ground to a fine powder 65 Similar compositions may also be granulated to form water soluble granules (SG).

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Wettable powders (WP) may be prepared by mixing a compound of the invention with one or more solid diluents or carriers, one or more wetting agents and, preferably, one or more dispersing agents and, optionally, one or more suspending agents to facilitate the dispersion in liquids. The mixture is then ground to a fine powder. Similar compositions may also be granulated to form water dispersible granules (WG).

Granules (GR) may be formed either by granulating a mixture of a compound of the invention and one or more powdered solid diluents or carriers, or from pre-formed blank granules by absorbing a compound of the invention (or a solution thereof, in a suitable agent) in a porous granular material (such as pumice, attapulgite clays, fuller's earth, kieselguhr, diatomaceous earths or ground corn cobs) or by adsorbing a compound of the invention (or a solution thereof, in a suitable agent) on to a hard core material (such as sands, silicates, mineral carbonates, sulfates or phosphates) and drying if necessary. Agents which are commonly used to aid absorption or adsorption include solvents (such as aliphatic and aromatic petroleum solvents, alcohols, ethers, ketones and esters) and sticking agents (such as polyvinyl acetates, polyvinyl alcohols, dextrins, sugars and vegetable oils). One or more other additives may also be included in granules (for example an emulsifying agent, wetting agent or dispersing agent).

Dispersible Concentrates (DC) may be prepared by dissolving a compound of the invention in water or an organic solvent, such as a ketone, alcohol or glycol ether. These solutions may contain a surface active agent (for example to improve water dilution or prevent crystallization in a spray tank).

Emulsifiable concentrates (EC) or oil-in-water emulsions (EW) may be prepared by dissolving a compound of the invention in an organic solvent (optionally containing one or more wetting agents, one or more emulsifying agents or a mixture of said agents). Suitable organic solvents for use in ECs include aromatic hydrocarbons (such as alkylbenzenes or alkylnaphthalenes, exemplified by SOLVESSO 100, SOLVESSO 150 and SOLVESSO 200; SOLVESSO is a Registered Trade Mark), ketones (such as cyclohexanone or methylcyclohexanone) and alcohols (such as benzyl alcohol, furfuryl alcohol or butanol), N-alkylpyrrolidones (such as N-methylpyrrolidone or N-octylpyrrolidone), dimethyl amides of fatty acids (such as C₈-C₁₀ fatty acid dimethylamide) and chlorinated hydrocarbons. An EC product may spontaneously emulsify on addition to water, to produce an emulsion with sufficient stability to allow spray application through appropriate equipment. Preparation of an EW involves obtaining a compound of the invention either as a liquid (if it is not a liquid at room temperature, it may be melted at a reasonable temperature, typically below 70° C.) or in solution (by dissolving it in an appropriate solvent) and then emulsifying the resultant liquid or solution into water containing one or more SFAs, under high shear, to produce an emulsion. Suitable solvents for use in EWs include vegetable oils, chlorinated hydrocarbons (such as chlorobenzenes), aromatic solvents (such as alkylbenzenes or alkylnaphthalenes) and other appropriate organic solvents which have a low solubility in water.

Microemulsions (ME) may be prepared by mixing water with a blend of one or more solvents with one or more SFAs, to produce spontaneously a thermodynamically stable isotropic liquid formulation. A compound of the invention is present initially in either the water or the solvent/SFA blend. Suitable solvents for use in MEs include those hereinbefore described for use in ECs or in EWs. An ME may be either

an oil-in-water or a water-in-oil system (which system is present may be determined by conductivity measurements) and may be suitable for mixing water-soluble and oil-soluble pesticides in the same formulation. An ME is suitable for dilution into water, either remaining as a microemulsion or ⁵ forming a conventional oil-in-water emulsion.

Suspension concentrates (SC) may comprise aqueous or non-aqueous suspensions of finely divided insoluble solid particles of a compound of the invention. SCs may be prepared by ball or bead milling the solid compound of the invention in a suitable medium, optionally with one or more dispersing agents, to produce a fine particle suspension of the compound. One or more wetting agents may be included in the composition and a suspending agent may be included to reduce the rate at which the particles settle. Alternatively, a compound of the invention may be dry milled and added to water, containing agents hereinbefore described, to produce the desired end product.

Aerosol formulations comprise a compound of the invention and a suitable propellant (for example n-butane). A compound of the invention may also be dissolved or dispersed in a suitable medium (for example water or a water miscible liquid, such as n-propanol) to provide compositions for use in non-pressurized, hand-actuated spray pumps.

A compound of the invention may be mixed in the dry state with a pyrotechnic mixture to form a composition suitable for generating, in an enclosed space, a smoke containing the compound.

Capsule suspensions (CS) may be prepared in a manner 30 similar to the preparation of EW formulations but with an additional polymerization stage such that an aqueous dispersion of oil droplets is obtained, in which each oil droplet is encapsulated by a polymeric shell and contains a compound of the invention and, optionally, a carrier or diluent 35 therefor. The polymeric shell may be produced by either an interfacial polycondensation reaction or by a coacervation procedure. The compositions may provide for controlled release of the compound of the invention and they may be used for seed treatment. A compound of the invention may 40 also be formulated in a biodegradable polymeric matrix to provide a slow, controlled release of the compound.

A composition may include one or more additives to improve the biological performance of the composition (for example by improving wetting, retention or distribution on 45 surfaces; resistance to rain on treated surfaces; or uptake or mobility of a compound of the invention). Such additives include surface active agents, spray additives based on oils, for example certain mineral oils or natural plant oils (such as soy bean and rape seed oil), and blends of these with other 50 bio-enhancing adjuvants (ingredients which may aid or modify the action of a compound of the invention).

A compound of the invention may also be formulated for use as a seed treatment, for example as a powder composition, including a powder for dry seed treatment (DS), a water 55 soluble powder (SS) or a water dispersible powder for slurry treatment (WS), or as a liquid composition, including a flowable concentrate (FS), a solution (LS) or a capsule suspension (CS). The preparations of DS, SS, WS, FS and LS compositions are very similar to those of, respectively, 60 DP, SP, WP, SC and DC compositions described above. Compositions for treating seed may include an agent for assisting the adhesion of the composition to the seed (for example a mineral oil or a film-forming barrier).

Wetting agents, dispersing agents and emulsifying agents 65 may be surface SFAs of the cationic, anionic, amphoteric or non-ionic type.

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Suitable SFAs of the cationic type include quaternary ammonium compounds (for example cetyltrimethyl ammonium bromide), imidazolines and amine salts.

Suitable anionic SFAs include alkali metals salts of fatty acids, salts of aliphatic monoesters of sulfuric acid (for example sodium lauryl sulfate), salts of sulfonated aromatic compounds (for example sodium dodecylbenzenesulfonate, calcium dodecylbenzenesulfonate, butylnaphthalene sulfonate and mixtures of sodium di-isopropyl- and tri-isopropyl-naphthalene sulfonates), ether sulfates, alcohol ether sulfates (for example sodium laureth-3-sulfate), ether carboxylates (for example sodium laureth-3-carboxylate), phosphate esters (products from the reaction between one or more fatty alcohols and phosphoric acid (predominately mono-esters) or phosphorus pentoxide (predominately diesters), for example the reaction between lauryl alcohol and tetraphosphoric acid; additionally these products may be ethoxylated), sulfosuccinamates, paraffin or olefine sulfonates, taurates and lignosulfonates.

Suitable SFAs of the amphoteric type include betaines, propionates and glycinates.

Suitable SFAs of the non-ionic type include condensation products of alkylene oxides, such as ethylene oxide, propylene oxide, butylene oxide or mixtures thereof, with fatty alcohols (such as oleyl alcohol or cetyl alcohol) or with alkylphenols (such as octylphenol, nonylphenol or octylcresol); partial esters derived from long chain fatty acids or hexitol anhydrides; condensation products of said partial esters with ethylene oxide; block polymers (comprising ethylene oxide and propylene oxide); alkanolamides; simple esters (for example fatty acid polyethylene glycol esters); amine oxides (for example lauryl dimethyl amine oxide); and lecithins.

Suitable suspending agents include hydrophilic colloids (such as polysaccharides, polyvinylpyrrolidone or sodium carboxymethylcellulose) and swelling clays (such as bentonite or attapulgite).

A compound of the invention may be applied by any of the known means of applying pesticidal compounds. For example, it may be applied, formulated or unformulated, to the pests or to a locus of the pests (such as a habitat of the pests, or a growing plant liable to infestation by the pests) or to any part of the plant, including the foliage, stems, branches or roots, to the seed before it is planted or to other media in which plants are growing or are to be planted (such as soil surrounding the roots, the soil generally, paddy water or hydroponic culture systems), directly or it may be sprayed on, dusted on, applied by dipping, applied as a cream or paste formulation, applied as a vapor or applied through distribution or incorporation of a composition (such as a granular composition or a composition packed in a water-soluble bag) in soil or an aqueous environment.

A compound of the invention may also be injected into plants or sprayed onto vegetation using electrodynamic spraying techniques or other low volume methods, or applied by land or aerial irrigation systems.

Compositions for use as aqueous preparations (aqueous solutions or dispersions) are generally supplied in the form of a concentrate containing a high proportion of the active ingredient, the concentrate being added to water before use. These concentrates, which may include DCs, SCs, ECs, EWs, MEs, SGs, SPs, WPs, WGs and CSs, are often required to withstand storage for prolonged periods and, after such storage, to be capable of addition to water to form aqueous preparations which remain homogeneous for a sufficient time to enable them to be applied by conventional spray equipment. Such aqueous preparations may contain

varying amounts of a compound of the invention (for example 0.0001 to 10%, by weight) depending upon the purpose for which they are to be used.

The invention will now be illustrated by the following non-limiting Examples. All citations are incorporated by reference.

BIOLOGICAL EXAMPLES

TABLE A

(Ia)

$$F_3C$$
 X^1
 X^1
 X^2
 X^3
 X^4
 X^4

	X1	R1	R4	R5	chem- istry at *	chem- istry at **		Al
A1	3,5- dichloro-	methyl	СН3	Н	R/S	R	30	A
A2	phenyl 3,5- dichloro-	2,2,2- trifluoro-	СН3	Н	R/S	R	30	A
A3	phenyl 3,5- dichloro-	ethyl ethyl	СН3	Н	S	R	35	Αź
A4	phenyl 3,5- dichloro- phenyl	ethyl	СН3	Н	R/S	S		A2
A5	3,5- dichloro- phenyl	methyl	СН3	Н	R/S	S	40	A2
A 6	3,4,5- trichloro- phenyl	ethyl	СН3	Н	R/S	R		A2
A7	3,5- dichloro- 4-bromo-	ethyl	CH3	Н	R/S	R	45	A2
A8	phenyl 3,5- dichloro- 4-fluoro-	ethyl	СН3	Н	R/S	R		
A9	phenyl 3,5- dichloro- phenyl	2,2,2- trifluoro- ethyl	СН3	Н	S	R	50	A2
A10	3,5- trifluoro- methyl- 4-chloro-	ethyl	СН3	Н	R/S	R	55	A2
A11	phenyl 3-chloro- 5-fluoro- phenyl	ethyl	СН3	Н	R/S	R		A2
A12	3,5- dichloro- phenyl	2,2- difluoro- ethyl	СН3	Н	S	R	60	A2
A13	3,4,5- trichloro- phenyl	2,2,2- trifluoro- ethyl	СН3	Н	S	R		A3
A14	3,5- dichloro- 4-fluoro- phenyl	2,2,2- trifluoro- ethyl	СНЗ	Н	S	R	65	A 3

$$F_{3}C$$

$$X^{l}$$

$$R^{5}$$

$$R^{4}$$

$$R^{4}$$

$$R^{4}$$

$$R^{4}$$

$$R^{4}$$

$$R^{1}$$

15		X1	R1	R4	R5	Stereo- chem- istry at *	Stereo- chem- istry at **
	A15	3-chloro- 5-tri- fluoro-	ethyl	СН3	Н	S	R
20		methyl- phenyl					
	A16	3-chloro- 5-bromo- phenyl	ethyl	СН3	Н	S	R
25	A17	3-chloro- 5-bromo- phenyl	2,2,2- trifluoro- ethyl	СН3	Н	S	R
	A18	3,5- dichloro- phenyl	ethyl	СН3	Н	S	R/S
30	A19	3,5- dichloro- phenyl	2,2,2- trifluoro- ethyl	СН3	Н	S	R/S
	A20	3,5- dichloro- phenyl	2,2- difluoro- ethyl	СН3	Н	S	R/S
35	A21	3,5- dichloro- 4-fluoro- phenyl	Ethyl	СН3	Н	S	R/S
	A22	3,5- dichloro- phenyl	methyl	СН3	Н	S	R/S
40	A23	3,4,5- trichloro- phenyl	2,2,2- trifluoro- ethyl	СН3	Н	S	R/S
	A24	3,5- dichloro- 4-fluoro-	ethyl	СН3	Н	S	R
45	A25	phenyl 3,5- dichloro- 4-fluoro-	ethyl	Cl	Н	S	R
50	A26	phenyl 3-chloro, 5-tri- fluoro-	ethyl	СН—СН—	СН—СН	S	R
	A27	methyl- phenyl 3,5- dichloro-	ethyl	Br	Н	S	R
55	A28	phenyl 3,5- dichloro-	ethyl	CF3	Н	S	R
	A29	phenyl 3,4,5- trichloro-	ethyl	CF3	Н	S	R
60	A30	phenyl 3-tri fluoro- methyl- phenyl	ethyl	СНЗ	Н	S	R
65	A31	3,4,5- trichloro- phenyl	ethyl	CF3	Н	S	R

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50

$$F_3C$$
 X^1
 X^1
 X^2
 X^3
 X^4
 X^4

	X1	R1	R4	R5	Stereo- chem- istry at *	Stereo- chem- istry at **
A32	3,5- dichloro- phenyl	ethyl	CF3	Н	S	R
A33	3,4,5- trichloro- phenyl	ethyl	Br	Н	S	R
A34	3,5- dichloro- phenyl	ethyl	Br	Н	S	R
A35	3,4,5- trichloro- phenyl	ethyl	СН3	Н	S	R

Table A provides compounds of formula (Ia) wherein X^1 and R^1 have the definitions shown below.

TABLE B

$$F_3C \xrightarrow{\hspace*{1cm} N \hspace*{1cm} } \overset{R^5}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad \qquad }} \overset{(Ib)}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad \qquad }} \overset{R^4}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad \qquad }} \overset{(Ib)}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad }} \overset{R^4}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad \qquad }} \overset{(Ib)}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad }} \overset{R^4}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad }} \overset{(Ib)}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad }} \overset{R^4}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad }} \overset{(Ib)}{\underset{\hspace*{1cm} N \hspace*{1cm} }{ \qquad }$$

	X1	R1	R4	R5	Stereo- chemistry at *	Stereo- chemistry at **	55
E	3,5-dichloro- phenyl	ethyl	СН3	Н	R/S	R	
E	3,5-dichloro- phenyl	ethyl	СН3	Η	R	R	60
E	33 3,5-dichloro- phenyl	2,2,2- trifluoroethyl	СН3	Н	R	R	

Table B provides compounds of formula (Ib) wherein X^1 and R^1 have the definitions shown below.

$$F_3C$$

$$X^1$$

$$X^1$$

$$X^2$$

$$X^3$$

$$X^4$$

	X1	R1	R4	R5	Stereo- chemistry at *	Stereo- chemistry at **
C1	3,5-dichloro- phenyl	ethyl	СН3	Н	R/S	R

R/S indicates a racemic mixture.

Table C provides compounds of formula (Ic) wherein X^1 and R^1 have the definitions shown below.

Nephotettix virescens (Green Leafhopper)

 25 Rice seedlings are treated with the diluted test solutions in a spray chamber. After drying, plants are infested with 10 $\rm N_{\rm 5}$ nymphs (3 replicates). 5 days after the treatment samples are checked for mortality.

Compound	Test	Application rate/ppm	Control/%
A 9	Nephotettix virescens (Green	200	100
	leafhopper)	50	100
		12	100
		3	100
A3	Nephotettix virescens (Green	200	100
	leafhopper)	50	100
	* * *	12	100
		3	90

Nilaparvata lugens (Brown Plant Hopper)

Rice seedlings are treated with the diluted test solutions in a spray chamber. After drying, plants are infested with $20\,\mathrm{N}_3$ nymphs (2 replicates). 6-12 days after the treatment samples are checked for mortality

Compound	Rate/ppm	Control/%
A1	50	98
	25	93
	12.5	93
A2	50	99
	25	98
	12.5	80
A3	50	100
	25	100
	12.5	100
A4	50	100
	25	93
	12.5	88
A5	50	98
	25	65
	12.5	0
A6	50	100
	25	100
	12.5	100
A7	50	100
	25	100
	12.5	100

A25

A26

A27

A28

A29

A30

12.5

12.5

25

12.5

12.5

12.5

12.5

12.5

	33 -continued		> , . , . , - ,		34 -continued	
Compound	Rate/ppm	Control/%		Compound	Rate/ppm	Control/%
A8	50 25	100 100	5	A31	50	100
A9	12.5 50 25	100 100 100			25 12.5	88 55
A10	12.5 50	98 100	10	A32	50 25	95 50
	25 12.5	100 100		A33	12.5 50	0 98
A11	50 25 12.5	100 55 80			25 12.5	98 80
A12	50 25	100 100	15	A34	50 25	100 100
A13	12.5 50 25	100 98 100		A35	12.5 50	98 98
A14	12.5 50	75 100	20		25 12.5	100 93
A15	25 12.5 50	100 100 100		B1	50 25	90 78
AD	25 12.5	99 99	25	B2	12.5 50	0 88
A16	50 25	98 97			25 12.5	70 15
A17	12.5 50 25	95 100 100	30	В3	50 25	100 100
A18	12.5 50	100 100		C1	12.5 50	100 95
A19	25 12.5 50	100 100 100			25 12.5	15
	25 12.5	100 100	35		12.5	
A20	50 25	100 100		ilo suppressalis (I		
A21	12.5 50 25	100 100 100	⁴⁰ cut	and transferred i	nto a petri dish o	drying the leaves are coated with wet filter
A22	12.5 50 25	100 100 100	a f	ilter tissue and a	plastic lid. 5 day	e dish is covered with
A23	12.5 50	98 100	per 45 tra		is assessed (3 re	eplicates per concen-
A24	25 12.5 50	98 90 98	_			Application
	25	0.5	C	ompound Test		rate/ppm Control/%

	Compound	Test	Application rate/ppm	Control/%
50	A9	Chilo suppressalis (Rice stem borer)	0.8 0.2	100 100
	A3	Chilo suppressalis (Rice stem borer)	0.05 0.8 0.2 0.05	43 100 100 97

COMPARATIVE EXAMPLES

Compounds are tested according to the above methods. The results show that the compounds of the invention are 65 significantly more active against the indicated rice pests than structurally similar compounds, particularly at low rates of application.

COMPARATIVE TABLE 1

Compound of the invention
$\begin{array}{c} Cl \\ \\ \\ Cl \\ \\ \\ Cl \\ \\ \\ \\ \\ \\ \\ \\ \\ $

Reference compound

$$\begin{array}{c}
CI \\
CI \\
CI
\end{array}$$

$$\begin{array}{c}
F_3C \\
O \\
N \\
O \\
\end{array}$$

$$\begin{array}{c}
I5 \\
N \\
N \\
O \\
\end{array}$$

$$\begin{array}{c}
20 \\
\end{array}$$

Compound Test		Application rate/ ppm	Control/ %	25
Compound of the invention Reference	Chilo suppressalis (Rice stem borer) Chilo suppressalis	0.8 0.2 0.8	100 53 43	
compound	(Rice stem borer)	0.8	23	30

COMPARATIVE TABLE 2

Compound of the invention

Reference compound

$$\begin{array}{c} CI \\ \\ CI \\ \\ CI \\ \end{array}$$

Compound	Test	Appli- cation rate/ ppm	Control/
Compound of	Nilaparvata lugens	50	99
the invention	(Brown plant hopper)	25	98
		12.5	80
Reference	Nilaparvata lugens	50	45
compound	(Brown plant hopper)	25	0
		12.5	0

The compound of the invention and reference compound are compounds B5 and B4 respectively from WO 2011/067272.

The invention claimed is:

1. A method for controlling the infestation of at least one of stemborer, leaffolder, hopper, Gall midge, whorl maggot, Rice bug, and Black bug in rice, the method comprising applying to a crop of rice plants, the locus thereof, or propagation material thereof, a compound of formula I

wherein
$$-B^1-B^2-B^3$$
— is $-C=N-O-$, $-C=N-CH_2-$, or $-C=CH_2-O-$;

L is a direct bond or methylene;

A¹ and A² are C—H, or one of A¹ and A² is C—H and the other is N;

X¹ is group X

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$$\begin{array}{c} \text{H} \\ \\ \text{(R}^3)_p \end{array}$$

 R^1 is C_1 - C_4 alkyl, C_1 - C_4 haloalkyl or C_3 - C_6 cycloalkyl;

R² is chlorodifluoromethyl or trifluoromethyl;

each R³ is independently bromo, chloro, fluoro or trifluoromethyl;

 R^4 is hydrogen, halogen, methyl, halomethyl or cyano; R^5 is hydrogen;

or R⁴ and R⁵ together form a bridging 1,3-butadiene group;

p is 2 or 3.

2. The method according to claim 1, wherein the method is a method of controlling infestation of stemborer in rice.

- 3. A method of controlling infestation of a pest chosen from at least one of a stemborer and a leaffolder in a crop of useful plants comprising applying to a crop of useful plants,
 60 the locus thereof, or propagation material thereof, a compound of formula I as defined in claim 1.
 - 4. The method according to claim 1, wherein the method is a method of controlling infestation of leaffolder.
- 5. The method according to claim 1, wherein the method is a method of controlling infestation of a pest chosen from at least two of a hopper, a gallmidge, a whorl maggot, a Rice bug, and a Black bug.

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(I*)

6. A method of controlling infestation of a pest chosen from at least one of a hopper, a gallmidge, a whorl maggot, a Rice bug, and a Black bug in a crop of useful plants comprising applying to a crop of useful plants, the locus thereof, or propagation material thereof, a compound of formula I as defined in claim **1.**

7. The method according to claim 1, wherein A^1 and A^2 are C—H, R^2 is trifluoromethyl, and R^5 is hydrogen, wherein the method is a method of controlling infestation of $_{10}$ hoppers or stemborer.

8. The method according to claim **1**, wherein A¹ and A² are C—H, R² is trifluoromethyl, R⁴ is methyl, R⁵ is hydrogen, each R³ is chlorine, p is 2, wherein the method is a method of controlling infestation of hoppers or stemborer.

9. The method according to claim 1, wherein R^1 is methyl, ethyl, propyl, butyl, cyclopropyl, cyclobutyl, trifluoroethyl, difluoroethyl, wherein the method is a method of controlling infestation of hoppers or stemborer.

10. The method according to claim 1, wherein R^1 is ethyl or trifluoroethyl, wherein the method is a method of controlling infestation of hoppers or stemborer.

11. The method according to claim 1, wherein X^1 is 3,5-dichloro-phenyl, 3,5-dichloro-4-fluorophenyl-, 3,4,5-trichlorophenyl-, or 3,5-bis(trifluoromethyl)phenyl, wherein the method is a method of controlling infestation of hoppers or stemborer.

12. The method according to claim 1, wherein $-B^1$ — $_{30}$ B^2 — B^3 —is -C—N—O—, wherein the method is a method of controlling infestation of hoppers or stemborer.

13. The method according to claim 1, wherein the compound of formula I is a mixture of compounds I* and I**

-continued (I**)

wherein the molar proportion of compound 1** compared to the total amount of both enantiomers is greater than 50%, wherein the method is a method of controlling infestation of hoppers or stemborer.

14. A method according to claim 1, wherein the compound of formula I is a mixture of compounds I' and I"

(S = S stereochemistry, R = R stereochemistry)

wherein the molar proportion of compound I" compared to the total amount of both enantiomers is greater than 50%, wherein the method is a method of controlling infestation of hoppers or stemborer.

* * * * *